ABOUT

inVentiv Health Clinical is recognized as a leading provider of global product development services to pharmaceutical, biotechnology, generic drug, and medical device companies, offering therapeutically specialized capabilities for Phase I-IV clinical development, bioanalysis, and clinical staffing from a single professional to an entire functional team. For intelligent solutions needed to accelerate high quality product development programs of all sizes around the world, inVentiv Health Clinical can help you transform promising ideas into commercial reality.

CAPABILITIES

Phase I-IIa

For proof-of-concept, First in Human (FIH) or bioequivalency studies, our extensive database of study participants and relationships with leading hospitals ensure rapid recruitment for clinical studies. In addition to a dedicated project manager for each study, we deploy a team of experts and specialists in quality assurance, and scientific and regulatory affairs to custom-fit a program to your needs.

Phase IIb-III

inVentiv Health Clinical combines extensive therapeutic knowledge with a commitment to quality and proven operational expertise to meet our clients' product development goals. Our dedicated project teams approach their work with a deep understanding of the specific requirements of each therapeutic category. We offer expertise and services for clinical trial design and a full range of clinical trial services, including biostatistics, clinical monitoring, data management, global safety and pharmacovigilance, regulatory consulting, medical writing, project management, and full-service patient recruitment and retention.

Exclusive access to UnitedHealth Care claims data allows us to have real-time, robust insight into data from more than 90,000 de-identified patients in the US to help accelerate recruitment. We will consult with you to apply the data and develop a protocol to set a strong foundation for an efficient, safe and cost-effective study. Our global footprint and therapeutic expertise can help you progress drugs and medical devices closer to regulatory submission, anywhere in the world.

Late Stage

inVentiv Health Clinical has a dedicated team of experts in strategic and operational planning, observational studies and patient registries, health economics and outcomes research, safety/risk management and epidemiology, and traditional interventional studies to guide clients through the post-approval environment.

Strategic Resourcing

inVentiv Health Clinical offers robust capabilities with a complete range of functional and clinical staffing services. Our broad capabilities and worldwide resources allow us to tailor resourcing solutions on a global scale. We deliver the right resources, customized to our clients' needs, and continually look for ways to reduce costs, enhance quality, and deliver timely results.

Drawing upon the combined experience of more than 3,000 employees and over 200,000 professional candidates, inVentiv Health Clinical is uniquely qualified to help its clients address the growing pressures to enhance innovation, expedite pipelines and improve efficiencies, while lowering costs. The company currently has 32 active partnerships dating back to 1999.

Bioanalytical

inVentiv Health Clinical offers bioanalytical support through our two state-of-the-art GLP-compliant bioanalytical laboratories, an extensive list of validated assays, knowledgeable scientists, and skilled technicians. Our experts develop, optimize, and validate analytical methods and rapidly process sample analysis of drugs from toxicokinetic, PK, bioavailability, bioequivalence, and all stages of clinical studies for both small and large molecules.

Technology

inVentiv Health Clinical provides comprehensive software support services for a variety of clinical trial data management and EDC systems. Our technology services are supported by fully validated business continuity plans, redundant data storage and timely backup procedures to ensure the integrity of your data. In addition, we offer a complete IVRS/IWRS and eDiary solution that is easy to deploy, scalable and 21 CFR Part 11 compliant.

AREAS OF THERAPEUTIC EXPERTISE

- Oncology
- Neuroscience
- Cardiovascular
- Endocrinology/Metabolism
- Rheumatology
- Pain Management
- Infectious Diseases/Vaccines
- Respiratory/Pulmonology
- Dermatology
- Ophthalmology
- Urology
- Nephrology
- Women's Health